



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

APR 28 2004

Food and Drug Administration  
9200 Corporate Boulevard  
Rockville MD 20850

Mr. Terence Lim  
Associate Manager, Quality Assurance Regulatory Affairs  
WRP Asia Pacific Sdn Bhd  
Lot 1, Jalan 3, Kawasan Perusahaan  
Bandar Baru Salak Tinggi  
43900 Sepang, Selangor Darul Ehsan  
MALAYSIA

Re: K040645  
Trade/Device Name: Dermagrip-G and Multiple Customers Trade name Green Color,  
Aloe Vera Coated Latex Examination Gloves With Proteen Content Labeling Claim  
(50 Micrograms or Less Water Extractable Protein)  
Regulation Number: 880.6250  
Regulation Name: Patient Examination Glove  
Regulatory Class: I  
Product Code: LYY  
Dated: March 9, 2004  
Received: March 11, 2004

Dear Mr. Lim:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (301) 594-4618. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>

Sincerely yours,



Chiu Lin, Ph.D.

Director

Division of Anesthesiology, General Hospital,  
Infection Control and Dental Devices  
Office of Device Evaluation  
Center for Devices and  
Radiological Health

Enclosure



WRP Asia Pacific Sdn Bhd

CA 141792

## Indications for Use

510(k) Number (if known): \_\_\_\_\_

Device Name: DERMAGRIE-G AND MULTIPLE CUSTOMER TRADE NAME  
GREEN COLOR, ALOE VERA COATED, LATEX PATIENT  
EXAMINATION GLOVES WITH PROTEIN CONTENT  
LABELING CLAIM (50 MICROGRAMS OR LESS WATER  
EXTRACTABLE PROTEIN)

### Indications For Use

A patient examination glove is a disposable device intended for medical purposes that is worn on the examiner's hands or finger to prevent contamination between patient and examiner.  
(21 CFR 880.6250)

Prescription Use \_\_\_\_\_  
(Part 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use \_\_\_\_\_  
(21 CFR 807 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF  
NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Kevin Muley

(Division Sign-Off)  
Division of Anesthesiology, General Hospital,  
Infection Control, Dental Devices

510(k) Number: K040645

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